

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
19 September 2002 (19.09.2002)

PCT

(10) International Publication Number
WO 02/072746 A1

(51) International Patent Classification⁷: C11D 3/39, 3/28

(21) International Application Number: PCT/EP02/01811

(22) International Filing Date: 18 February 2002 (18.02.2002)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0106285.0 14 March 2001 (14.03.2001) GB
01305810.2 5 July 2001 (05.07.2001) EP

(71) Applicant (for AE, AG, AU, BB, BZ, CA, CY, GB, GD, GH, GM, IE, IL, KE, LC, LK, LS, MN, MW, NZ, OM, SD, SG, SI, SZ, TT, TZ, UG, ZA, ZM, ZW only): UNILEVER PLC [GB/GB]; Unilever House, Blackfriars, London EC4P 4BQ (GB).

(71) Applicant (for all designated States except AE, AG, AU, BB, BZ, CA, CY, GB, GD, GH, GM, IE, IL, IN, KE, LC, LK, LS, MN, MW, NZ, OM, SD, SG, SI, SZ, TT, TZ, UG, ZA, ZM, ZW): UNILEVER NV [NL/NL]; Weena NV, NL-3013 AL Rotterdam (NL).

(71) Applicant (for IN only): HINDUSTAN LEVER LIMITED [IN/IN]; Hindustan Lever House, 165 / 166 Backbay Reclamation, Maharashtra, 400 020 Mumbai (IN).

(72) Inventors: HAGE, Ronald; Unilever R & D Vlaardingen, Olivier van Noortlaan 120, NL-3133 AT Vlaardingen (NL).

HERMANT, Roelant, Mathijs; Unilever R & D Vlaardingen, Olivier van Noortlaan 120, NL-3133 AT Vlaardingen (NL). VEERMAN, Simon, Marinus; Unilever R & D Vlaardingen, Olivier van Noortlaan 120, NL-3133 AT Vlaardingen (NL).

(74) Agents: ELLIOTT, Peter, William et al.; Unilever PLC, Patent Department, Colworth House, Sharnbrook, Bedford, Bedfordshire MK44 1LQ (GB).

(81) Designated States (national): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:
— with international search report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

WO 02/072746 A1

(54) Title: BLEACHING CATALYSTS WITH UNSATURATED SURFACTANT AND ANTIOXIDANTS

(57) Abstract: The invention relates to catalytically bleaching substrates, especially laundry fabrics, with a bleaching catalyst in the presence of an unsaturated organic surfactant and at least two antioxidants as activity moderator.

BLEACHING CATALYSTS WITH UNSATURATED SURFACTANT AND ANTIOXIDANTS

FIELD OF INVENTION

This invention relates to the protection of unsaturated
5 moieties in a bleaching composition.

BACKGROUND OF INVENTION

The use of bleaching catalysts for stain removal has been
developed over recent years. The recent discovery that some
10 catalysts are capable of bleaching effectively in the
absence of an added peroxy source has recently become the
focus of some interest, for example: WO9965905; WO0012667;
WO0012808; WO0029537, and, WO0060045.

15 UK patent application 0030877.5, filed 18-Dec-2000,
discloses the use of an unsaturated surfactant as a bleach
enhancement catalyst. However, there are stability problems
associated with the interaction of the unsaturated
surfactant and bleach enhancement catalyst.

20

DETAILED DESCRIPTION OF THE INVENTION

We have found that in some instances an unsaturated
surfactant is degraded by an air bleaching catalyst in a
non-desirable way. We have also found that in some
25 instances a peroxy bleaching catalyst together with a
peroxy species degrades an unsaturated surfactant in a non-
desirable way. A solution to this problem is provided by the
presence of an antioxidant, the presence of which still
permits air bleaching of stains.

30

- 2 -

It is an object of the present invention to provide a composition that reduces the problem of malodour. This is provided by the presence of an antioxidant. It is a further object to provide the composition that has a reduced level of antioxidant present. This is provided by the presence of a combination of at least two antioxidants in the composition.

The combination of the at least two antioxidants providing in a solution containing oleic acid an effective reduction in the formation of hexanal from the oleic acid under ambient atmospheric conditions by a factor of at least three in comparison with same composition having a molar equivalent of a single antioxidant equivalent to the combined molar concentration of the at least two antioxidants, said single antioxidant being one of the at least two antioxidants.

The present invention provides a bleaching composition comprising an organic ligand which forms a complex with a transition metal for bleaching a substrate with a group selected from:

- a) atmospheric oxygen, the bleaching composition upon addition to an aqueous medium providing an aqueous bleaching medium substantially devoid of a peroxygen bleach or a peroxy-based or peroxy-generating bleach system; and,
 - b) a peroxygen bleach or source thereof,
- together with a surfactant having an allylic hydrogen, said surfactant having an HLB of greater than 2, and at least two antioxidants, whereby the combination of the at least two antioxidants provides in a solution containing oleic acid an

- 3 -

effective reduction in the formation of hexanal from the oleic acid under ambient atmospheric conditions by a factor of at least three in comparison with same bleaching composition having a molar equivalent of a single
5 antioxidant equivalent to the combined molar concentration of the at least two antioxidants, said single antioxidant being one of the at least two antioxidants.

In a preferred embodiment of the present invention is
10 provided a bleaching composition for bleaching a substrate, the bleaching composition comprising:

- (i) an organic ligand which forms a complex with a transition metal for bleaching with oxygen sourced from the air;
- 15 (ii) 0.01 to 60 wt/wt% of a surfactant having an HLB of greater than 15, the surfactant a sodium salt of an unsaturated carboxylic acid having an allylic hydrogen; and,
- (iii) 0.001 to 5% wt/wt% of at least two antioxidants in a
20 molar ratio of at least 5%,

said bleaching composition comprising less than 2% mMol of peroxide per Kg, wherein upon addition of the bleaching composition to an aqueous solution and in the presence of the substrate and least 10 % of any bleaching of the
25 substrate is effected by oxygen sourced from the air and wherein the combination of the at least two antioxidants provides in a solution containing oleic acid an effective reduction in the formation of hexanal from the oleic acid under ambient atmospheric conditions by a factor of at least
30 three in comparison with same bleaching composition having a molar equivalent of a single antioxidant equivalent to the

- 4 -

combined molar concentration of the at least two antioxidants, said single antioxidant being one of the at least two antioxidants.

- 5 It is preferred that the bleaching composition is substantially devoid of a peroxygen bleach or a peroxy-based or peroxy-generating bleach system. Nevertheless, as another aspect of the present invention a peroxy source may be present such that "air bleaching" is suppressed.
- 10 Generally, "air bleaching" catalysts are capable of operating in a peroxy bleaching mode.

The surfactant having an allylic hydrogen has an HLB (hydrophilic/lipophilic balance) greater than 2, more preferably greater than 5, and most preferably greater than 10. Ideally, if the surfactant is a charged species the HLB is greater than 15. For a discussion of HLB the reader is directed to an article by Griffin, W. C. in J. Soc. Cosmetic Chemists Vol. 1 page 311, 1945 and Davies, J. T. and Rideal, E. K. in Interfacial Phenomena, Acad. Press, NY, 1961, pages 371 to 382. The HLB value requirement reflects the importance of the rate of solubility and dispersibility of the surfactant having an allylic hydrogen from the bleaching composition to the aqueous wash medium in conjunction with surface activity towards the substrate being washed. The threshold value of HLB as required excludes compounds that have an allylic which do not have the required surfactant properties, for example linoleic or oleic acid have an HLB of 0.8.

- 5 -

It is preferred that the surfactant having an allylic hydrogen has a CMC of 2×10^{-2} M or less. It is most preferred that the surfactant is anionic has a critical micelle concentration value of 3×10^{-3} M or less. Generally, a surfactant will form a micelle when present in an aqueous solution above a specific concentration that is intrinsic to the surfactant. A micelle is a neutral or electrically charged colloidal particle, consisting of oriented molecules. Above what is known as the *critical micelle concentration* CMC amphiphilic compounds tend to adopt specific aggregates in aqueous solution. The tendency is to avoid contact between their hydrophobic alkyl chains and the aqueous environment and to form an internal hydrophobic phase. Such compounds can form monomolecular layers [monolayers] at the air-water boundary and bimolecular layers [bilayers] between two aqueous compartments. Micelles are spherically closed monolayers. This CMC criterion is another aspect that aids reduction of catalyst deposit.

The property required is that the surfactant used in the present invention is and forms a micelle at a concentration of 2×10^{-2} M and below in an aqueous solution at a temperature of 25°C. One skilled in the art will be aware that the standard CMC is measured in deionized water and that the presence of other components in solution, e.g. surfactants or ions in solution will perturb the CMC value. The CMC values and requirement thereof as described herein are measured under standard conditions (N. M. Van Os, J. R. Haak, and L. A. M. Rupert, *Physico Chemical Properties of Selected Anionic Cationic and Nonionic Surfactants* Elsevier 1993; Kresheck, G. C. *Surfactants-In water a comparative*

- 6 -

treatise- (ed. F. Franks) Chapter 2 pp 95-197 Plenum Press
1971, New York; and, Mukerjee, P. and Mysels K. J. Critical
Micelle Concentrations of Aqueous Surfactant Systems, NSRDS-
NBS 36, National Bureau of Standards. US Gov. Print office
5 1971, Washington, DC).

The present invention has particular utility as a bleaching
composition in a commercial "air bleaching" liquid and
granular "air bleaching" or peroxy bleach format. The
10 degradation of unsaturated components during storage in the
absence of an antioxidant often results in the formation of
mal odour components due to the degradation of unsaturated
compounds. The composition also serves to reduce the
degradation of unsaturated compounds during the wash.

15 The composition of the present invention, in an air
bleaching mode, is preferably substantially devoid of a
peroxy bleach or a peroxy-based or peroxy-generating
bleach system. The term "substantially devoid of a peroxygen
20 bleach or a peroxy-based or peroxy-generating bleach
system" should be construed within spirit of the invention.
It is preferred that the composition has as low a content of
a peroxy species present as possible. Nevertheless,
autoxidation is something that is very difficult to avoid
25 and as a result small levels of peroxy species may be
present. These small levels may be as high as 2% but are
preferably below 2%. The level of peroxide present is
expressed in mMol of hydroperoxide (-OOH) present per Kg.
The additionally added organic compounds having labile CH's,
30 for example allylic, benzylic, -C(O)H, and -CRH-O-R', are
particularly susceptible to autoxidation and hence may

- 7 -

contribute more to this level of peroxy species than other components. However the presence of an antioxidant in the composition will likely serve to reduce the presence of adventitious peroxy species by reducing chain reactions.

- 5 The composition of the present invention bleaches a substrate with at least 10 %, preferably at least 50 % and optimally at least 90 % of any bleaching of the substrate being effected by oxygen sourced from the air.
- 10 When only a peroxyacid is present as a peroxy bleaching species in a bleaching medium with a bleaching catalyst $[\text{total peroxy present}] = [\text{RC(O)OOH}] + [\text{RC(O)OO}^-]$. When a mixture of hydrogen peroxide and peroxyacid are present in this medium $[\text{total peroxy present}] = [\text{RC(O)OOH}] + [\text{RC(O)OO}^-]$
- 15 $+ [\text{H}_2\text{O}_2] + [\text{HOO}^-]$. In some instances, the peroxy species will be relatively unreactive and hence the dominant conditions for "air bleaching" will be still be met by a relatively high level of peroxy species present. The different peroxy species will react at different rates with
- 20 an "air bleaching catalyst" but what is essential, for "air bleaching mode" is that $k[\text{air cat}][\text{peroxy}]$ is sufficiently small that $k[\text{air cat}][\text{O}_2]$ dominates to the extent that at least 10 % of any bleaching of the substrate is effected by oxygen sourced from the air when the composition is for use
- 25 in an air bleaching mode. When the composition is used in a peroxy mode there is sufficient peroxy species present to dominate and suppress "air bleaching" in the medium.

- The present invention extends to a commercial package
- 30 comprising the bleaching composition according to the present invention together with instructions for its use.

- 8 -

Any suitable textile that is susceptible to bleaching or one that one might wish to subject to bleaching may be used. Preferably the textile is a laundry fabric or garment.

- 5 In a preferred embodiment, the method according to the present invention is carried out on a laundry fabric using an aqueous treatment liquor. In particular, the treatment may be effected in a wash cycle for cleaning laundry. More preferably, the treatment is carried out in an aqueous
10 detergent bleach wash liquid, preferably in a washing machine.

- The composition of the present invention whilst providing an improved amount protection to unsaturated compounds permits
15 a bleaching activity of at least 25 %, preferably at least 50 %, equivalent to same composition devoid of antioxidant.

- A unit dose as used herein is a particular amount of the bleaching composition used for a type of wash. The unit
20 dose may be in the form of a defined volume of powder, granules or tablet.

Antioxidant

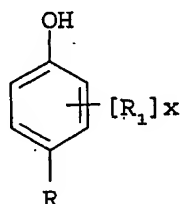
- The compositions of the present invention will comprise an
25 effective amount of the anti-oxidant, preferably from about 0.001 % more preferably from about 0.1%, most preferably from about 0.2% to about 10%, preferably to about 5%, more preferably to about 1% by weight of an anti-oxidant. Anti-oxidants are substances as described in Kirk-Othmers (Vol 3,
30 pg 424) and in Uhlmans Encyclopedia (Vol 3, pg 91).

- 9 -

It is preferred that the at least two antioxidants are present in the composition in a molar ratio of at least 5%, preferably at least 10%, most preferably 25% (0.1:0.025-see experimental).

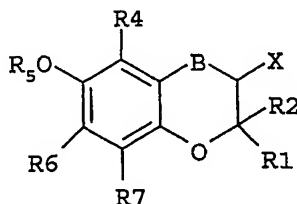
5

One class of anti-oxidants suitable for use in the present invention is alkylated phenols having the general formula:



wherein R is C1-C22 linear or branched alkyl, preferably methyl or branched C3-C6 alkyl; C3-C6 alkoxy, preferably methoxy; R1 is a C3-C6 branched alkyl, preferably tert-butyl; x is 1 or 2. Hindered phenolic compounds are preferred as antioxidant.

15 Another class of anti-oxidants suitable for use in the present invention is a benzofuran or benzopyran derivative having the formula:



wherein R1 and R2 are each independently alkyl or R1 and R2 can be taken together to form a C5-C6 cyclic hydrocarbyl moiety; B is absent or CH2; R4 is C1-C6 alkyl; R5 is hydrogen or -C(O)R3 wherein R3 is hydrogen or C1-C19 alkyl; R6 is C1-C6 alkyl; R7 is hydrogen or C1-C6 alkyl; X is -

20

- 10 -

CH₂OH, or - CH₂A wherein A is a nitrogen comprising unit, phenyl, or substituted phenyl. Preferred nitrogen comprising A units include amino, pyrrolidino, piperidino, morpholino, piperazino, and mixtures thereof.

5

Other suitable antioxidants are found as follows. A derivative of α -tocopherol, 6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid (Trolox[™]).

Anti-oxidants/radical scavengers such as ascorbic acid
10 (vitamin C) and its salts, tocopherol (vitamin E), tocopherol sorbate, other esters of tocopherol, butylated hydroxy benzoic acids and their salts, gallic acid and its alkyl esters, especially propyl gallate, uric acid and its salts and alkyl esters, sorbic acid and its salts, the
15 ascorbyl esters of fatty acids, amines (e.g., N,N-diethylhydroxylamine, amino-guanidine), sulfhydryl compounds (e.g., glutathione), and dihydroxy fumaric acid and its salts may be used.

20 Non-limiting examples of anti-oxidants suitable for use in the present invention include phenols *inter alia* 2,6-di-tert-butylphenol, 2,6-di-tert-butyl-4-methylphenol, mixtures of 2 and 3-tert-butyl-4-methoxyphenol, and other ingredients including include propyl gallate, tert-
25 butylhydroquinone, benzoic acid derivatives such as methoxy benzoic acid, methylbenzoic acid, dichloro benzoic acid, dimethyl benzoic acid, 5-hydroxy-2,2,4,6,7-pentamethyl-2,3-dihydro-1-benzofuran-3-one, 5-hydroxy-3-methylene-2,2,4,6,7-pentamethyl-2,3-dihydro-benzofuran, 5-
30 benzyloxy-3-hydroxymethyl-2,2,4,6,7-pentamethyl-2,3-dihydro-1-benzofuran, 3-hydroxymethyl-5-methoxy-2,2,4,6,7-

- 11 -

pentamethyl-2,3-dihydro-1-benzofuran, vitamin C (ascorbic acid), and Ethoxyquine (1,2-dihydro-6-ethoxy-2,2,4-trimethylchinolin) marketed under the name Raluquin[™] by the company Raschig[™].

5

Preferred radical scavengers for use herein include di-tert-butyl hydroxy toluene (BHT), α -tocopherol, hydroquinone, 2,2,4-trimethyl-1,2-dihydroquinoline, di-tert-butyl hydroquinone, mono-tert-butyl hydroquinone, tert-butyl-
10 hydroxy anisole, benzoic acid and derivatives thereof, like alkoxylated benzoic acids, as for example, trimethoxy benzoic acid (TMBA), toluic acid, catechol, t-butyl catechol, benzylamine, 1,1,3-tris(2-methyl-4-hydroxy-5-t-butylphenyl) butane, N-propyl-gallate or mixtures thereof
15 and highly preferred is di-tert-butyl hydroxy toluene.

Surfactant Having an Allylic Hydrogen

To benefit from the enhancement of bleaching activity it is preferred that the surfactant having an allylic hydrogen is
20 present in the composition such that a unit dose provides at least 0.01 g/l, more preferably at least 0.5g /l, and most preferably at least 0.1g /l, concentration of the unsaturated organic compound in a wash. The surfactant having an allylic hydrogen may be present in the composition
25 in the range of 0.01 to 60%, preferably 0.1 to 20% and most preferably 10% w/w.

There are many classes of surfactants having an allylic hydrogen that will work with the present invention to
30 enhance air bleaching. As one skilled in the art is aware a surfactant having an allylic hydrogen (enhancer) may be

- 12 -

found in: neutral species, and charged species, i.e., cationic species, anionic species, and zwitterionic species. It is preferred that the surfactant having an allylic hydrogen contains a hydrophilic group thereby providing the organic compound unassociated or as a micelle in an aqueous medium. It also is preferred that the surfactant having an allylic hydrogen is provided in the form of an alkali metal salt, preferably sodium, of an unsaturated carboxylic acid.

One skilled in the art will appreciate that benzene and toluene are considered unsaturated but neither possess allylic hydrogens per se. The homolytic bond dissociation energy (BDE) for benzene (C_6H_5-H) is 110.9 kcal/mol (298 K) makes benzene moieties per se unsuitable to promote enhanced bleaching. The surfactant used to enhance bleaching according to the present invention has a hydrogen atom covalently bound to an alpha-carbon that is alpha to a Sp^2 - Sp^2 hybridized bond (other than Sp^2 - Sp^2 hybridized bonds found in a cyclic aromatic system) e.g., as shown as underlined in the following formula $CH_2=CH-\underline{CH_2}-CH_3$. It is most preferred that the surfactant having an allylic hydrogen has a molecular weight of at least 80 and a bond dissociation energy of less than 95 kcal/mol, most preferably below 90 kcal/mol, and even more preferably below 85 kcal/mol. Below is a table of bond strengths (298 K) obtained from: The handbook of Chemistry and Physics 73rd edition, CRC Press.

- 13 -

Compound	BDE ΔH (kcal/mol)
(CH ₃) ₃ CH	93.3 \pm 0.5
H-CH ₂ OCH ₃)	93 \pm 1
C ₆ H ₅ -H	110.9 \pm 2.0
H-CMe ₂ OH	91 \pm 1
CH ₃ CH ₃	100.3 \pm 1
CH ₂ =CH-CH ₂ -CH ₃	83.1 \pm 2.2
CH ₂ =CH-CH ₃	86.3 \pm 1.5
C ₆ H ₅ -CH ₃	88.0 \pm 1
CH ₃ CH=CHCH=CH ₂	83 \pm 3

1) Unsaturated Soap (Unsaturated Anionic Surfactant)

5 The unsaturated fatty acid soap used preferably contains from about 16 to about 22 carbon atoms, preferably in a straight chain configuration. Preferably the number of carbon atoms in the unsaturated fatty acid soap is from about 16 to about 18.

10

This unsaturated soap, in common with other anionic detergents and other anionic materials in the detergent compositions of this invention, has a cation, which renders the soap water-soluble and/or dispersible. Suitable cations include sodium, potassium, ammonium, monoethanolammonium, diethanolammonium, triethanolammonium, tetramethylammonium, etc. cations. Sodium ions are preferred although in liquid formulations potassium, monoethanolammonium,

15

- 14 -

diethanolammonium, and triethanolammonium cations are useful.

The unsaturated soaps are made from natural oils that often
5 contain one or more unsaturated groups and consist of mixtures of components. It is clear that hydrolysis of these natural components yield mixtures of soaps, of which at least one of the components contain one or more unsaturated groups. Examples of natural oils are sunflower
10 oil, olive oil, cottonseed oil, linseed oil, safflower oil, sesame oil, palm oil, corn oil, peanut oil, soybean oil, castor oil, coconut oil, canola oil, cod liver oil and the like, that give mixtures of soaps of which at least one of them has at least one unsaturated group. However, also
15 hydrolysis products of purified oils, as listed above, may be employed. Other examples of soaps include those derived from erucic acid,

2) Unsaturated Surfactant (Unsaturated Cationic)

20 As one skilled in the art will appreciate such an unsaturated cationic may be manufactured, for example, by adding an unsaturated alkyl halide to an amine thus forming an unsaturated cationic.

25 In principle the cationic surfactants exhibit the same requirements as listed above for the unsaturated soap materials, except they need to be quarternised. Without limiting the scope of the invention, suitable cationics may be formed by preparing the quaternary salts from alcohols
30 that were obtained from the corresponding fatty acid (as defined under 1; from oils containing unsaturated bonds).

- 15 -

Examples of cationic surfactants based on natural oils include oleylbis(2-hydroxyethyl)methylammonium chloride and ditallow fatty alkyl dimethyl ammonium chloride.

5 3) Unsaturated Neutral Surfactant

An example of a non-ionic (neutral) surfactant is found in alkoxyated non-ionic surfactants. In common with the ionic surfactants as described above the surfactant has an allylic hydrogen.

10

Bleach Catalyst

The bleach catalyst per se may be selected from a wide range of organic molecules (ligands) and complexes thereof.

Suitable organic molecules (ligands) and complexes for use

15 with the present invention are found, for example in:

GB 9906474.3; GB 9907714.1; GB 98309168.7, GB 98309169.5;

GB 9027415.0 and GB 9907713.3; DE 19755493; EP 999050;

WO-A-9534628; EP-A-458379; EP 0909809; United States Patent

4,728,455; WO-A-98/39098; WO-A-98/39406, WO 9748787,

20 WO 0029537; WO 0052124, and WO0060045 the complexes and

organic molecule (ligand) precursors of which are herein

incorporated by reference. The air bleaching catalysts as

used herein should not be construed as an peroxy-generating

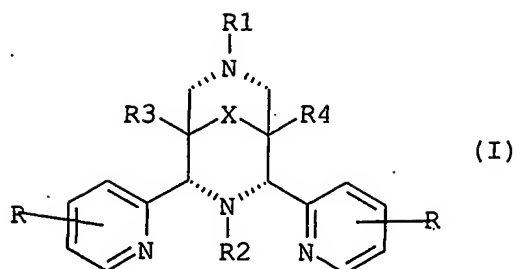
system, alone or in combination with other substrates,

25 irrespective of how they bleaching action works.

Another example of an air bleaching catalyst is a ligand or transition metal catalyst thereof of a ligand having the formula (I):

30

- 16 -



wherein each R is independently selected from: hydrogen, hydroxyl, and C1-C4-alkyl;

5 R1 and R2 are independently selected from:

C1-C4-alkyl,

C6-C10-aryl, and,

a group containing a heteroatom capable of coordinating to a transition metal, wherein at least one of R1 and R2 is the group containing the heteroatom;

10 R3 and R4 are independently selected from hydrogen, C1-C8 alkyl, C1-C8-alkyl-O-C1-C8-alkyl, C1-C8-alkyl-O-C6-C10-aryl, C6-C10-aryl, C1-C8-hydroxyalkyl, and $-(CH_2)_nC(O)OR_5$

wherein R5 is C1-C4-alkyl, n is from 0 to 4, and mixtures

15 thereof; and,

X is selected from $C=O$, $-[C(R_6)_2]_y-$ wherein Y is from 0 to 3 each R6 is independently selected from hydrogen, hydroxyl, C1-C4-alkoxy and C1-C4-alkyl.

20 It is preferred that the group containing the heteroatom is: a heterocycloalkyl: selected from the group consisting of: pyrrolinyl; pyrrolidinyl; morpholinyl; piperidinyl; piperazinyl; hexamethylene imine; 1,4-piperazinyl; tetrahydrothiophenyl; tetrahydrofuranlyl; tetrahydropyranylyl; and oxazolidinyl, wherein the heterocycloalkyl may be

25

- 17 -

- connected to the ligand via any atom in the ring of the selected heterocycloalkyl,
a -C1-C6-alkyl-heterocycloalkyl, wherein the heterocycloalkyl of the -C1-C6-heterocycloalkyl is selected
5 from the group consisting of: piperidinyl; piperidine; 1,4-piperazine, tetrahydrothiophene; tetrahydrofuran; pyrrolidine; and tetrahydropyran, wherein the heterocycloalkyl may be connected to the -C1-C6-alkyl via any atom in the ring of the selected heterocycloalkyl,
10 a -C1-C6-alkyl-heteroaryl, wherein the heteroaryl of the -C1-C6-alkylheteroaryl is selected from the group consisting of: pyridinyl; pyrimidinyl; pyrazinyl; triazolyl; pyridazinyl; 1,3,5-triazinyl; quinolinyl; isoquinolinyl; quinoxalinyl; imidazolyl; pyrazolyl; benzimidazolyl;
15 thiazolyl; oxazolidinyl; pyrrolyl; carbazolyl; indolyl; and isoindolyl, wherein the heteroaryl may be connected to the -C1-C6-alkyl via any atom in the ring of the selected heteroaryl and the selected heteroaryl is optionally substituted by -C1-C4-alkyl,
20 a -C0-C6-alkyl-phenol or thiophenol,
a -C2-C4-alkyl-thiol, thioether or alcohol,
a -C2-C4-alkyl-amine, and
a -C2-C4-alkyl-carboxylate.
- 25 The ligand forms a complex with one or more transition metals, in the latter case for example as a dinuclear complex. Suitable transition metals include for example: manganese in oxidation states II-V, iron II-V, copper I-III, cobalt I-III, titanium II-IV, tungsten IV-VI, vanadium II-V
30 and molybdenum II-VI.

- 18 -

The transition metal complex preferably is of the general formula (AI):



5

in which:

M represents a metal selected from Mn(II)-(III)-(IV)-(V), Cu(I)-(II)-(III), Fe(II)-(III)-(IV)-(V), Co(I)-(II)-(III), Ti(II)-(III)-(IV), V(II)-(III)-(IV)-(V), Mo(II)-(III)-(IV)-(V)-(VI) and W(IV)-(V)-(VI), preferably from Fe(II)-(III)-(IV)-(V);

10

L represents the ligand, preferably N,N-bis(pyridin-2-yl-methyl)-1,1-bis(pyridin-2-yl)-1-aminoethane, or its protonated or deprotonated analogue;

15

X represents a coordinating species selected from any mono, bi or tri charged anions and any neutral molecules able to coordinate the metal in a mono, bi or tridentate manner;

Y represents any non-coordinated counter ion;

20 a represents an integer from 1 to 10;

k represents an integer from 1 to 10;

n represents zero or an integer from 1 to 10;

m represents zero or an integer from 1 to 20.

25 It is preferred that the organic molecule (ligand) or transition metal complex is present in the composition such that a unit dose provides at least 0.1 μ M of the organic molecule or transition metal complex thereof.

30 The present invention may be used in a peroxy bleaching mode in contrast to an air bleaching mode in which the

- 19 -

composition is substantially devoid of a peroxy source. However it is preferred to use the present invention in an air bleaching mode. In this instance a purely peroxy bleaching catalyst may be employed in contrast to an "air
5 bleaching" catalyst.

Peroxygen Bleach or Source Thereof

In a peroxy bleaching mode the composition of the present invention uses a peroxy species to bleach a substrate. The
10 peroxy bleaching species may be a compound which is capable of yielding hydrogen peroxide in aqueous solution. Hydrogen peroxide sources are well known in the art. They include the alkali metal peroxides, organic peroxides such as urea
15 perborates, percarbonates, perphosphates persilicates and persulphates. Mixtures of two or more such compounds may also be suitable.

Particularly preferred are sodium perborate tetrahydrate
20 and, especially, sodium perborate monohydrate. Sodium perborate monohydrate is preferred because of its high active oxygen content. Sodium percarbonate may also be preferred for environmental reasons. The amount thereof in the composition of the invention usually will be within the
25 range of about 1-35% by weight, preferably from 5-25% by weight. One skilled in the art will appreciate that these amounts may be reduced in the presence of a bleach precursor e.g., N,N,N',N'-tetraacetyl ethylene diamine (TAED).

30 Another suitable hydrogen peroxide generating system is a combination of a C1-C4 alkanol oxidase and a C1-C4 alkanol,

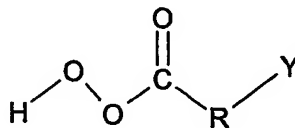
- 20 -

especially a combination of methanol oxidase (MOX) and ethanol. Such combinations are disclosed in International Application PCT/EP 94/03003 (Unilever), which is incorporated herein by reference.

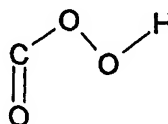
5

Alkylhydroxy peroxides are another class of peroxy bleaching compounds. Examples of these materials include cumene hydroperoxide and t-butyl hydroperoxide.

- 10 Organic peroxyacids may also be suitable as the peroxy bleaching compound. Such materials normally have the general formula:



- wherein R is an alkylene or substituted alkylene group
15 containing from 1 to about 20 carbon atoms, optionally having an internal amide linkage; or a phenylene or substituted phenylene group; and Y is hydrogen, halogen, alkyl, aryl, an imido-aromatic or non-aromatic group, a COOH
or



20

group or a quaternary ammonium group.

Typical monoperoxy acids useful herein include, for example:

- 25 (i) peroxybenzoic acid and ring-substituted peroxybenzoic acids, e.g. peroxy- α -naphthoic acid;

- 21 -

(ii) aliphatic, substituted aliphatic and arylalkyl monoperoxyacids, e.g. peroxyauric acid, peroxysearic acid and N,N-phthaloylamino peroxy caproic acid (PAP); and

5 (iii) 6-octylamino-6-oxo-peroxyhexanoic acid.

Typical diperoxyacids useful herein include, for example:

(iv) 1,12-diperoxydodecanedioic acid (DPDA);

10

(v) 1,9-diperoxyazelaic acid;

(vi) diperoxybrassicilic acid; diperoxysebasic acid and diperoxyisophthalic acid;

15

(vii) 2-decyldiperoxybutane-1,4-dioic acid; and

(viii) 4,4'-sulphonylbis peroxybenzoic acid.

20 Also inorganic peroxyacid compounds are suitable, such as for example potassium monopersulphate (MPS). If organic or inorganic peroxyacids are used as the peroxygen compound, the amount thereof will normally be within the range of about 2-10% by weight, preferably from 4-8% by weight.

25

Peroxyacid bleach precursors are known and amply described in literature, such as in the British Patents 836988; 864,798; 907,356; 1,003,310 and 1,519,351; German Patent 3,337,921; EP-A-0185522; EP-A-0174132; EP-A-0120591; and
30 U.S. Pat. Nos. 1,246,339; 3,332,882; 4,128,494; 4,412,934 and 4,675,393.

- 22 -

Another useful class of peroxyacid bleach precursors is that of the cationic i.e. quaternary ammonium substituted peroxyacid precursors as disclosed in US Pat. Nos. 4,751,015 and 4,397,757, in EP-A0284292 and EP-A-331,229. Examples of
5 peroxyacid bleach precursors of this class are:

2-(N,N,N-trimethyl ammonium) ethyl sodium-4-sulphonphenyl carbonate chloride (SPCC);

10 N-octyl-N,N-dimethyl-N10-carbophenoxy decyl ammonium chloride (ODC);

3-(N,N,N-trimethyl ammonium) propyl sodium-4-sulphophenyl carboxylate; and

15 N,N,N-trimethyl ammonium toluoyloxy benzene sulphonate.

A further special class of bleach precursors is formed by the cationic nitriles as disclosed in EP-A-303,520 and in
20 European Patent Specification No.'s 458,396 and 464,880.

Any one of these peroxyacid bleach precursors can be used in the present invention, though some may be more preferred than others.

25 Of the above classes of bleach precursors, the preferred classes are the esters, including acyl phenol sulphonates and acyl alkyl phenol sulphonates; the acyl-amides; and the quaternary ammonium substituted peroxyacid precursors
30 including the cationic nitriles.

- 23 -

Examples of said preferred peroxyacid bleach precursors or activators are sodium-4-benzoyloxy benzene sulphonate (SBOBS); N,N,N'N'-tetraacetyl ethylene diamine (TAED); sodium-1-methyl-2-benzoyloxy benzene-4-sulphonate; sodium-4-methyl-3-benzoyloxy benzoate; SPCC; trimethyl ammonium toluyloxy-benzene sulphonate; sodium nonanoyloxybenzene sulphonate (SNOBS); sodium 3,5,5-trimethyl hexanoyloxybenzene sulphonate (STHOBBS); and the substituted cationic nitriles.

10

Other classes of bleach precursors for use with the present invention are found in WO0015750, for example 6-(nonanamidocaproyl)oxybenzene sulphonate.

15 The precursors may be used in an amount of up to 12%, preferably from 2-10% by weight, of the composition.

The Detergent Composition.

The air bleach catalyst and unsaturated organic compound may be used in a detergent composition specifically suited for stain bleaching purposes, and this constitutes a second aspect of the invention. To that extent, the composition comprises a surfactant and optionally other conventional detergent ingredients. The invention in its second aspect provides an enzymatic detergent composition which comprises from 0.1 - 50 % by weight, based on the total detergent composition, of one or more surfactants. This surfactant system may in turn comprise 0 - 95 % by weight of one or more anionic surfactants and 5 to 100 % by weight of one or more nonionic surfactants. The surfactant system may additionally contain amphoteric or zwitterionic detergent

20
25
30

- 24 -

compounds, but this is not normally desired owing to their relatively high cost. The enzymatic detergent composition according to the invention will generally be used as a dilution in water of about 0.05 to 2%.

5

In general, the nonionic and anionic surfactants of the surfactant system may be chosen from the surfactants described "Surface Active Agents" Vol. 1, by Schwartz & Perry, Interscience 1949, Vol. 2 by Schwartz, Perry & Berch,
10 Interscience 1958, in the current edition of "McCutcheon's Emulsifiers and Detergents" published by Manufacturing Confectioners Company or in "Tenside-Taschenbuch", H. Stache, 2nd Edn., Carl Hauser Verlag, 1981.

15 Suitable nonionic detergent compounds which may be used include, in particular, the reaction products of compounds having a hydrophobic group and a reactive hydrogen atom, for example, aliphatic alcohols, acids, amides or alkyl phenols with alkylene oxides, especially ethylene oxide either alone
20 or with propylene oxide. Specific nonionic detergent compounds are C₆-C₂₂ alkyl phenol-ethylene oxide condensates, generally 5 to 25 EO, i.e. 5 to 25 units of ethylene oxide per molecule, and the condensation products of aliphatic C₈-C₁₈ primary or secondary linear or branched alcohols with
25 ethylene oxide, generally 5 to 40 EO.

Suitable anionic detergent compounds which may be used are usually water-soluble alkali metal salts of organic sulphates and sulphonates having alkyl radicals containing
30 from about 8 to about 22 carbon atoms, the term alkyl being used to include the alkyl portion of higher acyl radicals.

- 25 -

Examples of suitable synthetic anionic detergent compounds are sodium and potassium alkyl sulphates, especially those obtained by sulphating higher C₈-C₁₈ alcohols, produced for example from tallow or coconut oil, sodium and potassium

5 alkyl C₉-C₂₀ benzene sulphonates, particularly sodium linear secondary alkyl C₁₀-C₁₅ benzene sulphonates; and sodium alkyl glyceryl ether sulphates, especially those ethers of the higher alcohols derived from tallow or coconut oil and synthetic alcohols derived from petroleum. The preferred

10 anionic detergent compounds are sodium C₁₁-C₁₅ alkyl benzene sulphonates and sodium C₁₂-C₁₈ alkyl sulphates. Also applicable are surfactants such as those described in EP-A-328 177 (Unilever), which show resistance to

15 salting-out, the alkyl polyglycoside surfactants described in EP-A-070 074, and alkyl monoglycosides.

Preferred surfactant systems are mixtures of anionic with nonionic detergent active materials, in particular the groups and examples of anionic and nonionic surfactants

20 pointed out in EP-A-346 995 (Unilever). Especially preferred is surfactant system that is a mixture of an alkali metal salt of a C₁₆-C₁₈ primary alcohol sulphate together with a C₁₂-C₁₅ primary alcohol 3-7 EO ethoxylate.

25 The nonionic detergent is preferably present in amounts greater than 10%, e.g. 25-90% by weight of the surfactant system. Anionic surfactants can be present for example in amounts in the range from about 5% to about 40% by weight of the surfactant system.

30

- 26 -

The detergent composition may take any suitable physical form, such as a powder, granular composition, tablets, a paste or an anhydrous gel.

5 Enzymes

The detergent compositions of the present invention may additionally comprise one or more enzymes, which provide cleaning performance, fabric care and/or sanitation benefits.

10

Said enzymes include oxidoreductases, transferases, hydrolases, lyases, isomerases and ligases. Suitable members of these enzyme classes are described in Enzyme nomenclature 1992: recommendations of the Nomenclature Committee of the International Union of Biochemistry and Molecular Biology on the nomenclature and classification of enzymes, 1992, ISBN 0-12-227165-3, Academic Press.

15

The composition may contain additional enzymes as found in WO 01/00768 A1 page 15, line 25 to page 19, line 29, the contents of which are herein incorporated by reference. Builders, polymers and other enzymes as optional ingredients may also be present as found in WO0060045.

20

25 Suitable detergency builders as optional ingredients may also be present as found in WO0034427.

The composition of the present invention may be used for laundry cleaning, hard surface cleaning (including cleaning of lavatories, kitchen work surfaces, floors, mechanical ware washing etc.). As is generally known in the art,

30

- 27 -

bleaching compositions are also employed in waste-water treatment, pulp bleaching during the manufacture of paper, leather manufacture, dye transfer inhibition, food processing, starch bleaching, sterilisation, whitening in
5 oral hygiene preparations and/or contact lens disinfection.

In the context of the present invention, bleaching should be understood as relating generally to the decolourisation of stains or of other materials attached to or associated with
10 a substrate. However, it is envisaged that the present invention can be applied where a requirement is the removal and/or neutralisation by an oxidative bleaching reaction of malodours or other undesirable components attached to or otherwise associated with a substrate. Furthermore, in the
15 context of the present invention bleaching is to be understood as being restricted to any bleaching mechanism or process that does not require the presence of light or activation by light.

20 The invention will now be further illustrated by way of the following non-limiting examples:

EXAMPLES

25 Example 1

Synthesis of [(MeN4Py)FeCl]Cl

The ligand N,N-bis(pyridin-2-yl-methyl)-1,1-bis(pyridin-2-yl)-1-aminoethane (MeN4py) was prepared as described in EP 0
30 909 809 A2. The synthesis of the iron complex, [(MeN4Py)FeCl]Cl, has been described elsewhere (WO 0116271.

- 28 -

Liquid formulation A was prepared with 0.03% of [Fe(MeN4py)Cl]Cl by adding 7.5 mgs of the solid material in 25 ml liquid formulation A and optionally the anti-oxidant mixtures were added (see table 1 for exact formulations).

- 5 The mixture was stirred vigorously for 10 min and the liquids were then stored at 38 °C.

The anti-oxidants employed were: Trolox (6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid, Raluquin (1,2-dihydro-6-ethoxy-2,2,4-trimethylchinolin, vitamine C, Vitamine E (α-tocopherol, and a mixture of 10% α, 45% δ- and 45% γ-tocopherol (denoted as tocopherol mix). The latter system was 70% pure, the values given in the table are corrected for this purity.

15

A SPME GC-MS headspace analysis on a HP 6890 mass spectrometer (E.I.) was performed and some of the products analysed (e.g., no perfume components) are listed in the table below, after 1-4 weeks storage at 38 °C (see table).

- 20 The intensities of the signals were integrated and the typical error in the determinations was around 5%. Further the bleach performance on tomato-oil stains was assessed by using the method described below. The bleach performance experiments were done immediately after mixing
25 the catalyst without or with anti-oxidant mixture.

Bottles tests were done (25 mL solution), each bottle containing two tomato stained cloths (4x4 cm).

- The cloths were washed for 30 min at 40 °C. The dosage of
30 formulation A was 5 g/l. The water hardness used was 24 °FH. After the wash, the cloths were rinsed with water and

- 29 -

subsequently dried, and the change in reflectance at 460 nm was measured immediately after drying on a Minolta CM-3700d spectrophotometer including a UV-Vis filter before and after treatment (denoted as $t=0$ in the table). Subsequently, the washed cloths were stored for 24 hrs in a dry dark cupboard at ambient conditions and the cloths were measured again (after-wash bleaching process), denoted as $t=1$ in the table. The difference in ΔR between both reflectance values gives a measure of the bleaching performance of the system on the stain, i.e. a higher ΔR value corresponds to an improved bleaching performance.

The results for bleaching performance are shown in table 1.

- 30 -

Table 1: Amounts of aldehydes detected by GC-MS and bleach results of the liquid detergent formulations containing catalyst and anti-oxidants after storing the liquids at 38 °C.

Experi- ment	Cat.	Antioxidant (% in formulation)	Weeks storage	Octanal	Heptanal	Hexanal	ΔR (t=0)	ΔR (t=1)
1	-	-	4	658	294	113	15*	16*
2	+	-	4	3933	2870	2462	22*	38*
3	+	0.10 tocopherol mix	4	5907	3947	1551	20**	33**
4	+	1.04 tocopherol mix	4	912	682	450	18**	25**
5	+	0.025% Raluquin	1	3636	2752	3314	20***	26***
6	+	0.025% Trolox	1	3066	2253	3538	23***	32***
7	+	0.10 tocopherol mix + 0.025% Raluquin	4	806	663	443	15**	25**
8	+	0.10 tocopherol mix + 0.025% Trolox	4	823	698	479	16**	21**

5

* The bleach activity of the blanks (experiments 1 and 2) was determined after 3 weeks storage

** The bleach activity of experiments 3,4, 7, and 8) was determined after 4 weeks storage

10 *** The bleach activity of experiments 5 and 6) were determined after 1 week storage.

- 31 -

Composition formulation A:

Component	%
PAS	10%
Nonionic surfactant, ethoxylated fatty Alcohol type	18.4%
Oleic acid	10%
Deflocculating polymer, polymer A11 from EP346,995	1%
silicon oil to control foam	0.03%
KOH	4.1 %
NaOH	0.9%
Citric acid.H2O	5.5%
Glycerol	5%
Borax	1.9%
Anti-dye transfer polymer	0.3%
Protease	0.3%
Lipolase	0.37%
Amylase	0.15%
Perfume	0.47%

From the results presented in the table, one can draw the following conclusions:

5

1. Addition of the iron catalyst leads to an increased formation of various aldehydes (octanal, heptanal, hexanal) as detected by GC-MS. Without being bound to theory, one can infer that these products are most likely formed due to degradation of the unsaturated soap present under storage conditions. This assumption was tested by preparing a liquid containing a fully saturated soap added. No detectable amounts of these

10

- 32 -

aldehydes with the catalyst added were observed after 3 days storage.

2. Under the same conditions a clear signal of pyridyl-carboxaldehyde was observed (data not shown). Without being bound to theory, one can infer that this is most likely caused by decomposition of the iron catalyst during storage.
3. Addition of various anti-oxidants mixtures, leads to a dramatic improved stability upon storage as compared to the anti-oxidants as such, as detected by analysing the aldehyde formation. See experiments 7 and 8 in comparison to experiments 3-6.

15

Claims:

1. A bleaching composition comprising an organic ligand
5 which forms a complex with a transition metal for
bleaching a substrate with a group selected from:
 - a) atmospheric oxygen, the bleaching composition upon
addition to an aqueous medium providing an aqueous
10 bleaching medium substantially devoid of a peroxygen
bleach or a peroxy-based or peroxy-generating bleach
system; and,
 - b) a peroxygen bleach or source thereof,
- 15 together with a surfactant having an allylic hydrogen,
said surfactant having an HLB of greater than 2, and at
least two antioxidants, whereby the combination of the at
least two antioxidants provides in a solution containing
oleic acid an effective reduction in the formation of
20 hexanal from the oleic acid under ambient atmospheric
conditions by a factor of at least three in comparison
with same bleaching composition having a molar equivalent
of a single antioxidant equivalent to the combined molar
concentration of the at least two antioxidants, said
25 single antioxidant being one of the at least two
antioxidants.
2. A bleaching composition according claim 1, wherein
the antioxidants are selected from the group consisting
30 of: phenols and amines.
3. A bleaching composition according to claim 2,
wherein the antioxidants are hindered phenols.

4. A bleaching composition according to any preceding claim, wherein the mixture of antioxidants is present in the bleaching composition in the range from 0.001 to 5 wt %.

5. A bleaching composition according to claim 4, wherein the mixture of antioxidants is present in the bleaching composition in the range from 0.2 to 1 wt %.

6. A bleaching composition according to any preceding claim, wherein said at least two antioxidants are present in the composition in a molar ratio of at least 5%, preferably at least 10%, most preferably at least 25%.

7. A bleaching composition according to any preceding claim, wherein the antioxidants are selected from the group consisting of: di-tert-butyl hydroxy toluene, Ethoxyquine, α -tocopherol, and 6-hydroxy-2,5,7,8-tetra-methylchroman-2-carboxylic acid.

8. A bleaching composition according to any preceding claim, wherein the unsaturated compound has a hydrogen atom covalently bound to an alpha-carbon that is alpha to a Sp^2 - Sp^2 hybridized bond, said hydrogen having a homolytic bond dissociation energy of less than 95 kcal/mol, most preferably below 90 kcal/mol, and even more preferably below 85 kcal/mol.

9. A bleaching composition according to any preceding claim, wherein the surfactant is selected from the group of unsaturated neutral species.

- 35 -

10. A bleaching composition to any preceding claim,
wherein the surfactant is selected from the group of
unsaturated zwitterionic species.
- 5 11. A bleaching composition according to any preceding
claim, wherein the surfactant has an HLB of greater than
5.
12. A bleaching composition according to claim 11,
10 wherein the surfactant has an HLB of greater than 10.
13. A bleaching composition according to any preceding
claim, wherein the surfactant is present in the
composition in an amount such that a unit dose provides
15 at least 0.01 g/l concentration of the unsaturated
organic compound in a wash.
14. A bleaching composition according to any preceding
claim, wherein the surfactant has a CMC of 2×10^{-2} M or
20 less.
15. A bleaching composition according to claim 14,
wherein the surfactant is anionic and has a critical
micelle concentration value of 3×10^{-3} M or less.
25
16. A bleaching composition according to any preceding
claim, wherein the surfactant has molecular weight of at
least 80 and the allylic hydrogen has bond dissociation
energy of less than 90 kcal/mol.
30

- 36 -

17. A bleaching composition according to any preceding claim, wherein the surfactant is present in the composition in the range of 0.01 to 60 % wt/wt.
- 5 18. A bleaching composition according to claim 17, wherein the surfactant is present in the composition in the range of preferably 0.1 to 20% wt/wt.
- 10 19. A bleaching composition according to any preceding claim, wherein the composition comprises less than 0.1% of a peroxy source and at least 10 %, preferably at least 50 % and optimally at least 90 % of any bleaching of the substrate being effected by oxygen sourced from the air.
- 15 20. A bleaching composition according to claims 1 to 18, wherein the composition comprises a peroxy source.
- 20 21. A bleaching composition according to any preceding claim, wherein the organic substance is N,N-bis(pyridin-2-yl-methyl)-1,1-bis(pyridin-2-yl)-1-aminoethane.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 02/01811

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C11D3/39 C11D3/28

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C11D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 622 646 A (PRETTY ALASTAIR J ET AL) 22 April 1997 (1997-04-22) cited in the application column 1, line 43-50 column 10, line 18 -column 11, line 6 column 13, line 10-49 claims 1,9; example II	1-18,20, 21
A	WO 00 60045 A (PROCTER & GAMBLE (US)) 12 October 2000 (2000-10-12) cited in the application page 10, line 15 -page 14, line 5 page 36, line 18 -page 37, line 13 claim 7; example 2	1-21

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

7 June 2002

Date of mailing of the international search report

17/06/2002

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040. Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Pentek, E

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No
PCT/EP 02/01811

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
US 5622646	A	22-04-1997	AT 170553 T	15-09-1998
			CA 2187175 A1	19-10-1995
			DE 69504489 D1	08-10-1998
			DE 69504489 T2	20-05-1999
			EP 0754218 A1	22-01-1997
			JP 3009471 B2	14-02-2000
			JP 9511774 T	25-11-1997
			WO 9527774 A1	19-10-1995
<hr/>				
WO 0060045	A	12-10-2000	AU 4061900 A	23-10-2000
			WO 0060045 A1	12-10-2000
<hr/>				

THIS PAGE DELETED